

# EXHIBIT 1

**From:** Misbah Sherwani <MSherwani@actavis.com>  
**Sent:** Tuesday, April 15, 2008 2:50 PM  
**To:**  
**Subject:** Fw: List by Product  
**Attach:** 5-Sept 07 - Present Investigations by product.xls

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She's gone through both.....there should be no more surprises.  
----- Forwarded by Misbah Sherwani/QA/ELZ/Alpharma on 04/15/2008 02:48 PM -----

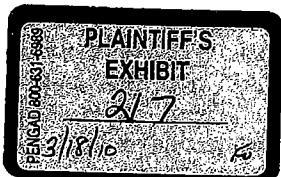
Michael Ponzo/Actavis@Exchange  
04/15/2008 02:17 PM

To  
Misbah Sherwani/QA/ELZ/Alpharma@Alpharma  
cc  
Anthony Castellazzo/Actavis@Exchange  
Subject  
List by Product

There were 2 that were incorrect (did not state stability OOS). Investigations 07-059 & 07-101.  
All stability OOS are now highlighted.

Mike

<<...>>



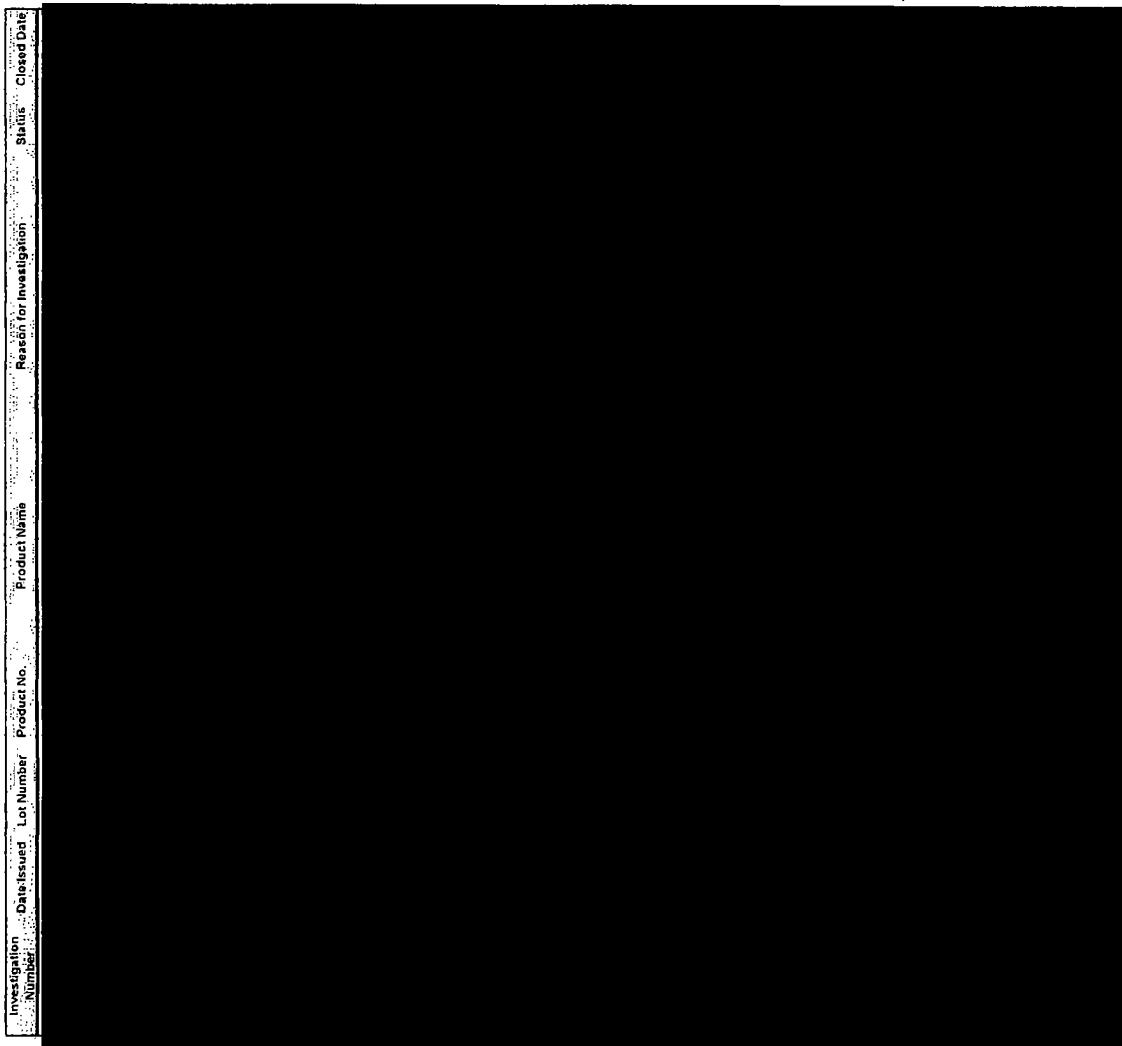
Investigation Date Issued: Lot Number Product Name Product Number Status Closed Date Reason for Investigation

Investigation Number	Date Issued	Lot Number	Product No.	Product Name	Reason for Investigation	Status	Closed Date
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Confidential - Subject Under Protective Order

ACT#V/000200006

Investigation Number	Date Issued	Lot Number	Product No.	Product Name	Reason for Investigation	Status	Closed Date
<b>Digitek (digoxin tablets, USP) 0.125 mg</b>							
07-093	12/05/07	70924A1	145	Digitek (digoxin tablets, USP) 0.125 mg	Two tablets of digoxin tablets 0.125 mg were found with approx. double the thickness from counter channels during packaging/filling operation.	closed	1/25/2008
08-011	1/29/2008	80044A1	145	Digitek (digoxin tablets, USP) 0.125 mg	One stainless steel screw was found in the tablet well of the BOSS-PACK filling machine during operation.	closed	3/3/2008
08-017	2/8/2008	80051A	145	Digitek (digoxin tablets, USP) 0.125 mg	During compression of tablets #2 and #6 the compression operator observed oil spots on some of the fractured tablets.	closed	3/6/2008
08-028	2/20/2008	80053A	145	Digitek (digoxin tablets, USP) 0.125 mg	Compression operator did not record the model detection test for the run on 2/15/08.	closed	2/23/2008
08-046	3/13/2008	7078A1	145	Digitek (digoxin tablets, USP) 0.125 mg	Zero stability testing was not conducted when CRT stability study was initiated.	open	
08-023	2/19/2008	80108A1	146	Digitek (digoxin tablets, USP) 0.25 mg	QA inspector & labeling operator verified an incorrect barcode chapter label.	closed	2/20/2008
08-029	2/20/2008	80152A	146	Digitek (digoxin tablets, USP) 0.25 mg	Product validation protocols were established for these batches and were not required. As a result additional samples were taken that were not required.	closed	3/14/2008
08-030	2/20/2008	80133A	146	Digitek (digoxin tablets, USP) 0.25 mg	Operator noticed tablets that were thinner than a typical tablet during the inspection of tablet #2.	closed	3/7/2008
08-031	2/25/2008	80152A	146	Digitek (digoxin tablets, USP) 0.25 mg	The compressed tablet description of NMPR # RV-1401 revision 00 is incorrect. The blend description in both the process validation and cleaning protocols is incorrect.	closed	3/13/2008
		80153A					
		80154A					
		80155A					



Investigation Number: Date Issued: Lot Number: Product No.: Reason for Investigation: Status: Closed Date:

ACTIV 00C299529

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